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(51) International Patent Classification 6: WO 98/22619 (11) International Publication Number: C12Q 1/37, C07D 235/20 **A1** (43) International Publication Date: 28 May 1998 (28.05.98) [CA/US]; 1019 Ashbury Street #3, San Francisco, CA PCT/US97/04450 (21) International Application Number: 94117 (US). MOORE, William, R. [US/US]; 1317 Balboa Avenue, Burlingame, CA 94010 (US). STROUD, Robert, 21 March 1997 (21.03.97) (22) International Filing Date: M. [GB/US]; 376 Castaneda Avenue, San Francisco, CA 94116 (US). (30) Priority Data: (74) Agents: DOW, Karen, B. et al.; Townsend and Townsend US 19 November 1996 (19.11.96) 08/746,986 and Crew LLP, 8th floor, Two Embarcadero Center, San Francisco, CA 94111-3834 (US). (63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application 08/746,986 (CON) (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, US BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, Filed on 19 November 1996 (19.11.96) HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, (71) Applicants (for all designated States except US): ARRIS PHARMACEUTICAL CORPORATION [US/US]; 385 US, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, Oyster Point Boulevard, South San Francisco, CA 94080 (US). THE REGENTS OF THE UNIVERSITY OF TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, CALIFORNIA [US/US]; 22nd floor, 300 Lakeside Drive, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, Oakland, CA 94612 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): CLARK, James, M. [US/US]; 2400 Kehoe Avenue, San Mateo, CA 94403 (US). Published ELROD, Kyle [US/US]; 4325 Sedge Street, Fremont, CA With international search report. 94555 (US). JENKINS, Thomas, E. [US/US]; 16 Redwood Drive, La Honda, CA 94020 (US). KATZ, Bradley, A.

(54) Title: METAL MEDIATED SERINE PROTEASE INHIBITORS

(57) Abstract

The present invention relates to serine protease inhibitors having a P site binding moiety and a divalent cation(s) chelating moiety which are pre-prepared as divalent cation(s) complexes or combined with the serine protease in the presence of divalent cation(s) and methods for identifying and using the inhibitors. The compounds are shown to have high inhibitory activity when complexed to divalent cation(s) and are useful in various processes associated with serine protease isolation and inhibition.

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